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=> s transferrin?(5a)receptor? 126255 TRANSFERRIN? 30355 RECEPTOR?

517 TRANSFERRIN?(5A)RECEPTOR?

=> s 11 and neisseria? 1439 NEISSERIA? 12 L1 AND NEISSERIA? 1.2

=> d 1-12

1. 5,708,149, Jan. 13, 1998, Method for producing purified

Haemophilus influenzae transferrin binding proteins; Sheena Loosmore, et

al., 530/418, 412, 413 :IMAGE AVAILABLE:

2. 5,698,438, Dec. 16, 1997, Bacterial hemoglobin receptor gene;

Stojiljkovic, et al., 435/252.3, 320.1; 536/23.7 :IMAGE AVAILABLE:

3. 5,693,463, Dec. 2, 1997, Method of ordering sequence binding preferences of a DNA-binding molecule; Cynthia A. Edwards, et al.,

7.23; 536/23.1; 935/76, 77 :IMAGE AVAILABLE:

4. 5,681,570, Oct. 28, 1997, Immunogenic conjugate molecules;

Yang, et al., 424/197.11, 203.1, 234.1, 244.1, 256.1; 514/54; 536/123.1

:IMAGE AVAILABLE:

5. 5,618,541, Apr. 8, 1997, Vaccine against **Neisseria** meningitidis

infections; Marie-Jose Quentin-Millet, 424/250.1, 249.1; 435/871 :IMAGE

AVAILABLE:

6. 5,618,540, Apr. 8, 1997, Subunit vaccine against **Neisseria** meningitidis infections and corresponding subunits in the purified

Marie J. Quentin-Millet, et al., 424/250.1, 249.1; 435/871 :IMAGE AVAILABLE:

7. 5,589,463, Dec. 31, 1996, Oral delivery of biologically active substances bound to vitamin B12; Gregory J. Russell-Jones, et al., 514/21; 424/194.1; 514/12, 52; 530/405; 536/26.4, 26.41, 26.44 :IMAGE AVAILABLE:

8. 5,578,444, Nov. 26, 1996, Sequence-directed DNA-binding

compositions and methods; Cynthia A. Edwards, et al., 435/6, 7.23; 536/23.1; 935/76, 77 :IMAGE AVAILABLE:

9. 5,428,023, Jun. 27, 1995, Oral delivery of biologically active substances bound to vitamin B12 or analogues thereof, Gregory J. Russell-Jones, et al., 514/21; 424/85.4, 193.1, 194.1; 514/2, 4, 6, 12,

15, 52; 530/303, 306, 313, 345, 351, 398, 399, 405, 409; 536/26.4, 26.41.

26.44 :IMAGE AVAILABLE:

10. 5,292,869, Mar. 8, 1994, Method for isolating and purifying **transferrin** and lactoferrin **receptor** proteins from bacteria

the preparation of vaccines containing the same; Anthony B. Schryvers,

530/413; 424/234.1, 249.1, 250.1, 255.1, 256.1; 530/350, 380, 394, 395,

400, 412, 417 :IMAGE AVAILABLE:

11. 5,141,743, Aug. 25, 1992, Method for isolating and purifying **transferrin** and lactoferrin **receptor** proteins and vaccines containing the same; Anthony B. Schryvers, 424/234.1, 249.1, 250.1,

255.1, 256.1, 278.1; 530/350, 395, 400, 413 :IMAGE AVAILABLE:

12. 4,943,522, Jul. 24, 1990, Lateral flow, non-bibulous membrane assav

protocols; Robert W. Eisinger, et al., 435/7.25; 422/55, 56, 57, 58, 101:

435/5, 7.21, 7.23, 7.32, 805, 810; 436/512, 514, 518, 520, 523,

807, 808, 810; D24/223 :IMAGE AVAILABLE:

=> d 1-12 date

L2: 1 of 12

TITLE: Method for producing purified recombinant Haemophilus

influenzae transferrin binding proteins

US PAT NO: 5,708,149 DATE ISSUED: Jan. 13, 1998 :IMAGE AVAILABLE:

APPL-NO: 08/487,890 DATE FILED: Jun. 7, 1995 REL-US-DATA: Continuation of Ser. No. 337,483, Nov. 8, 1994, which is a

> continuation-in-part of Ser. No. 175,116, Dec. 29, 1993, abandoned, which is a continuation-in-part of Ser. No. 148,968, Nov. 8, 1993, abandoned.

> > L2: 2 of 12

TITLE: Bacterial hemoglobin receptor gene

US PAT NO: 5,698,438 DATE ISSUED: Dec. 16,

1997

:IMAGE AVAILABLE:

APPL-NO: 08/326,670 DATE FILED: Oct. 18, 1994

L2: 3 of 12

TITLE: Method of ordering sequence binding preferences of a DNA-binding molecule

US PAT NO: DATE ISSUED: Dec. 2, 1997 5,693,463 :IMAGE AVAILABLE: DISCL-DATE: Apr. 26,

2011

APPL-NO: 07/996,783 DATE FILED: Dec. 23, 1992 REL-US-DATA: Continuation-in-part of Ser. No. 723,618, Jun. 27, 1991,

abandoned.

L2: 4 of 12

TITLE: Immunogenic conjugate molecules

US PAT NO: 5,681,570 1997

DATE ISSUED: Oct. 28,

:IMAGE AVAILABLE:

08/371,965 APPL-NO: DATE FILED: Jan. 12, 1995

L2: 5 of 12

TITLE: Vaccine against **Neisseria** meningitidis infections US PAT NO: 5,618,541 DATE ISSUED: Apr. 8, 1997

:IMAGE AVAILABLE:

APPL-NO: 08/066,167 DATE FILED: Jun. 2, 1993 FRN-PR. NO: 91 12177 FRN FILED: Oct. 3, 1991

FRN-PR. CO: France

PCT/FR92/00905 PCT-NO:

PCT-FILED: Sep. 29,

1992

Jun. 2, 1993 371-DATE:

102(E)-DATE: Jun. 2, 1993

PCT-PUB-NO: WO93/06861

PCT-PUB-DATE: Apr.

15, 1993

L2: 6 of 12

Subunit vaccine against **Neisseria** meningitidis TITLE: infections and corresponding subunits in the purified

state

US PAT NO: 5,618,540

DATE ISSUED: Apr. 8, 1997

:IMAGE AVAILABLE: APPL-NO:

08/064,174

DATE FILED: May 25, 1993

FRN-PR. NO: 91 12176 FRN-PR. CO: France

FRN FILED: Oct. 3, 1991

PCT-NO:

PCT-FILED: Sep. 29,

1992

PCT/FR92/00904

371-DATE: May 25, 1993 102(E)-DATE: May 25, 1993

PCT-PUB-NO: WO93/07172

PCT-PUB-DATE: Apr.

15, 1993

L2: 7 of 12

TITLE: Oral delivery of biologically active substances bound to vitamin B12

US PAT NO: 5,589,463

DATE ISSUED: Dec. 31,

1996

:IMAGE AVAILABLE:

APPL-NO: 08/479,635 DATE FILED: Jun. 7, 1995

FRN-PR. NO: PH2838

FRN FILED: Oct. 10, 1985

FRN-PR. CO: Australia

REL-US-DATA: Division of Ser. No. 61,343, May 17, 1993, Pat. No.

> 5,428,023, which is a continuation of Ser. No. 759,697, Sep. 9, 1991, abandoned, which is a continuation of Ser. No. 600,137, Oct. 19, 1990, abandoned, which is a continuation of Ser. No. 84,821, Jun. 9, 1987, abandoned.

> > L2: 8 of 12

Sequence-directed DNA-binding molecules TITLE: compositions and

methods

US PAT NO: 5,578,444

DATE ISSUED: Nov. 26,

1996

:IMAGE AVAILABLE:

APPL-NO: 08/171,389 DATE FILED: Dec. 20, 1993 REL-US-DATA: Continuation-in-part of Ser. No. 123,936, Sep. 17, 1993,

which is a continuation-in-part of Ser. No. 996,783, Dec. 23, 1992, which is a continuation-in-part of Ser. No. 723,618, Jun. 27, 1991, abandoned.

L2: 9 of 12

Oral delivery of biologically active substances bound to TITLE: vitamin B12 or analogues thereof

US PAT NO: 5,428,023

DATE ISSUED: Jun. 27,

1995

:IMAGE AVAILABLE:

PCT/AU86/00299

APPL-NO: 08/061,343 FRN-PR. NO: PH2838

DATE FILED: May 17, 1993

FRN-PR. CO: Australia

FRN FILED: Oct. 10, 1985 PCT-FILED: Oct. 10,

PCT-NO: 1986

Jun. 9, 1987 371-DATE:

102(E)-DATE: Jun. 9, 1987

PCT-PUB-NO: WO87/02351

PCT-PUB-DATE: Apr.

23, 1987

REL-US-DATA: Continuation of Ser. No. 759,697, Sep. 9, 1991, abandoned.

which is a continuation of Ser. No. 600,137, Oct. 19, 1990, abandoned, which is a continuation of Ser. No. 84,821, Jun. 9, 1987, abandoned.

L2: 10 of 12

Method for isolating and purifying **transferrin** and TITLE: lactoferrin **receptor** proteins from bacteria and the preparation of vaccines containing the same

US PAT NO: 5,292,869

DATE ISSUED: Mar. 8, 1994

:IMAGE AVAILABLE: 07/507,481

DATE FILED: Apr. 11, 1990

APPL-NO: REL-US-DATA: Continuation-in-part of Ser. No. 344,356, Apr. 27, 1989,

abandoned.

L2: 11 of 12

Method for isolating and purifying **transferrin** and TITLE: lactoferrin **receptor** proteins and vaccines

containing the same US PAT NO: 5,141,743

DATE ISSUED: Aug. 25,

:IMAGE AVAILABLE: APPL-NO: 07/639.365

DATE FILED: Jan. 10, 1991

REL-US-DATA: Continuation of Ser. No. 344,356, Apr. 27, 1989, abandoned.

L2: 12 of 12

Lateral flow, non-bibulous membrane assay protocols US PAT NO: 4,943,522 DATE ISSUED: Jul. 24, 1990

:IMAGE AVAILABLE:

DATE FILED: Aug. 10, 1988 APPL-NO: 07/230,642 REL-US-DATA: Continuation-in-part of Ser. No. 57,273, Jun. 1, 1987,

abandoned, and a continuation-in-part of Ser. No. 57,271, Jun. 1, 1987, abandoned.

=> d 5 ab

US PAT NO: 5,618,541 :IMAGE AVAILABLE:

12

ABSTRACT:

A vaccinal pharmaceutical composition which comprises, as therapeutic

agents, at least a first and a second molecule capable of binding to human transferrin; the said first molecule originating from a first strain of N. meningitidis which possesses a human **transferrin** **receptor** in which the lower molecular weight subunit (Tbp2) is recognised by an antiserum to the receptor of N. meningitidis strain 2394

(receptor 2394) and is not recognised by an antiserum to the receptor

N. meningitidis strain 2169 (receptor 2169); and at least a second molecule originating from a second strain of N. meningitidis which possesses a human **transferrin** **receptor** in which the lower molecular weight subunit (Tbp2) is recognised by an anti-receptor

antiserum and is not recognised by an anti-receptor 2394 antiserum.

=> d 6 ab

12

US PAT NO: 5.618.540 :IMAGE AVAILABLE:

L2: 6 of

1.2: 5 of

ABSTRACT:

The present invention relates to the lower molecular weight subunit

the human **transferrin** **receptor** of a strain of N. meningitidis, in

purified form, as well as to a vaccinal pharmaceutical composition intended for the prevention or attenuation of the effects of an N. meningitidis infection, containing the said subunit in purified form.

=> d 5 clms

US PAT NO: 5,618,541 :IMAGE AVAILABLE: 12

L2: 5 of

CLAIMS:

CLMS(1)

I claim:

1. A vaccinal pharmaceutical composition intended for preventing a **Neisseria** meningitidis infection, which comprises, as therapeutic agents, at least a first and a second molecule capable of binding to human transferrin which are either a human **transferrin**

receptor

of N. meningitidis or a subunit thereof; the said first molecule originating from a first strain of N. meningitidis which possesses a human **transferrin** **receptor** comprising a high molecular weight

subunit and a lower molecular weight subunit, and in which the lower

molecular weight subunit is recognised by an antiserum to the receptor of

N. meningitidis strain 2394 (receptor 2394) and is not recognised by an

antiserum to the receptor of N. meningitidis strain 2169 (receptor 2169);

and the said second molecule originating from a second strain of N. meningitidis which possesses a human **transferrin** **receptor** comprising a high molecular weight subunit and a lower molecular weight

subunit, and in which the lower molecular weight subunit is recognised by

an anti-receptor 2169 antiserum and is not recognised by an anti-receptor

2394 antiserum.

CLMS(2)

2. A vaccinal pharmaceutical composition according to claim 1, which

comprises, as therapeutic agents, at least a first and a second molecule

capable of binding human transferrin; the said first molecule originating

from a first strain of N. meningitidis which possesses a human
transferrin **receptor** in which the high molecular subunit
weight

and the lower molecular weight subunit are recognised by an antireceptor

2394 antiserum; and the said second molecule originating from a second

strain of N. meningitidis which possesses a human **transferrin**
receptor in which the high molecular weight subunit and the
lower

molecular weight subunit are recognised by an anti-receptor 2169 antiserum.

CLMS(3)

A vaccinal pharmaceutical composition according to claims 1 or

which comprises, as therapeutic agents, at least a first and a second

molecule capable of binding to human transferrin; the said first molecule

originating from a first strain of N. meningitidis which possesses a human **transferrin** **receptor** comprising a subunit of high molecular

weight of 100 kD approximately to 90 kD and a subunit of lower molecular $\,$

weight of 75 kD to 60 kD; and the said second molecule originating from a

second strain of N. meningitidis which possesses a human **transferrin**

receptor comprising of a subunit of high molecular weight of 100 kD

approximately to 90 kD and a subunit of lower molecular weight of 90 kD to 80 kD.

CLMS(4)

4. A vaccinal pharmaceutical composition according to claim 3, in which

the said first molecule originates from a first strain of N. meningitidis which possesses a human **transferrin** **receptor** comprising a subunit

of high molecular weight of 93-95 kD approximately and a subunit of lower molecular weight of 72 kD to 65 kD.

CLMS(5)

5. A vaccinal pharmaceutical composition according to claim 4, in

the said first molecule originates from a first strain of N. meningitidis which possesses a human **transferrin** **receptor** comprising a subunit

of high molecular weight of 93 kD approximately and a subunit of lower

molecular weight of 67-70 kD approximately.

CLMS(6)

6. A vaccinal pharmaceutical composition according to claim 5, in which

the said second molecule originates from a second strain of N. meningitidis which possesses a human **transferrin** **receptor** comprising a subunit of high molecular weight of 100 kD approximately to

95 kD and a subunit of lower molecular weight of 87 kD to 85 kD.

CLMS(7)

7. A vaccinal pharmaceutical composition according to claim 6, in which

the said second molecule originates from a second strain of N. meningitidis which possesses a human **transferrin** **receptor** comprising a subunit of high molecular weight of 98 kD approximately and

a subunit of lower molecular weight of 87 kD approximately.

CLMS(8)

8. A vaccinal pharmaceutical composition according to claim 1, in which

the said first and second molecules originate respectively from a first and second strain of N. meningitidis serogroup B.

CLMS(9)

9. A vaccinal pharmaceutical composition for preventing a **Neisseria**

meningitidis infection, which comprises therapeutic agents comprising a

first and a second molecule capable of binding to human transferrin which

are either a human **transferrin** **receptor** of N. meningitidis or a

subunit thereof; said first molecule originating from a first strain of N. meningitidis which possesses a human **transferrin**
receptor

comprising a high molecular weight subunit and a lower molecular weight

subunit, and in which the lower molecular weight subunit is recognized by

an antiserum to the receptor of N. meningitidis strain 2394 and is not recognized by an antiserum to the receptor of N. meningitidis strain 2169, and said second molecule, originating from a second strain of N.

meningitidis which possesses a human **transferrin** **receptor** comprising a high molecular weight subunit and a lower molecular weight

subunit, and in which the lower molecular weight subunit is recognized by

an antiserum to the receptor of N. meningitidis strain 2169 and is not recognized by an antiserum to the receptor of N. meningitidis strain 2394.

=> d 6 clms

US PAT NO: 5,618,540 :IMAGE AVAILABLE: L2: 6 of 12

CLAIMS:

CLMS(1)

We claim:

1. The lower molecular weight subunit of the human **transferrin**

receptor of a strain of N. meningitidis, in substantially purified form and in the absence of the higher molecular weight subunit of said
receptor.

CLMS(2)

2. The lower molecular weight subunit of the human **transferrin**
receptor of a strain of N. meningitidis serogroup B, in
substantially

purified form and in the absence of the higher molecular weight subunit of said receptor.

CLMS(3)

3. The lower molecular weight subunit of the human **transferrin**
receptor of a strain of N. meningitidis, in substantially purified form and in the absence of the higher molecular weight subunit of receptor, the said subunit having a molecular weight of 65 to 74 kD approximately.

CLMS(4)

4. The lower molecular weight subunit of the human **transferrin**
receptor of a strain of N. meningitidis strain 2394, in
substantially
purified form and in the absence of the higher molecular weight
subunit

CLMS(5)

of said receptor.

5. The lower molecular weight subunit of the human **transferrin**
receptor of a strain of N. meningitidis, in substantially purified
form and in the absence of the higher molecular weight subunit of
said

receptor; the said subunit having a molecular weight of 75 to 90 kD approximately.

CLMS(6)

The lower molecular weight subunit of the human **transferrin**
receptor of an N. meningitidis strain 2169, in substantially
purified

form and in the absence of the higher molecular weight subunit of said receptor.

CLMS(7)

 A vaccinal pharmaceutical composition which comprises, as therapeutic

agent, the lower molecular weight subunit of the human **transferrin**

receptor of at least one strain of N. meningitidis; in the absence of

the high molecular weight subunit of the said receptor.

CLMS(8)

8. A pharmaceutical composition according to claim 7, which comprises

the lower molecular weight subunit of the human **transferrin**
receptor of at least one strain of N. meningitidis serogroup B.

CLMS(9)

9. A pharmaceutical composition according to claim 7, which comprises,

as therapeutic agent, the lower molecular weight subunit of the human

transferring **receptor** of a strain of N. meningitidis; the said subunit having a molecular weight of 65 to 74 kD approximately.

CLMS(10)

 A pharmaceutical composition according to claim 9, which comprises,

as therapeutic agent, the lower molecular weight subunit of the human

transferrin **receptor** of N. meningitidis 2394.

CLMS(11)

11. A pharmaceutical composition according to claim 7, which comprises,

as therapeutic agent, the lower molecular weight subunit of the human

transferrin **receptor** of a strain of N. meningitidis; the said subunit having a molecular weight of 75 to 90 kD approximately.

CLMS(12)

A pharmaceutical composition according to claim 11, which comprises,

as therapeutic agent, the lower molecular weight subunit of the human

transferrin **receptor** of N. meningitidis 2169.

CLMS(13)

13. A pharmaceutical composition according to claim 7, which comprises,

as therapeutic agent:

- i) a first lower molecular weight subunit of the human
- **transferrin**
- **receptor** of a first strain of N. meningitidis; the said first subunit having a molecular weight of 65 to 74 kD approximately; and
- ii) a second lower molecular weight subunit of the human **transferrin**
- **receptor** of a second strain of N. meningitidis; the said second subunit having a molecular weight of 75 to 90 kD approximately; in the absence of the high molecular weight subunit of the said receptor

of the said first and second strains of N. meningitidis.

CLMS(14)

- 14. A pharmaceutical composition according to claim 13, which comprises,
- as therapeutic agent:
- i) the lower molecular weight subunit of the human **transferrin**
 receptor of N. meningitidis 2394; and
- ii) the lower molecular weight subunit of the human **transferrin**
 receptor of N. meningitidis 2169;
- in the absence of the high molecular weight subunit of the said receptor
- of N. meningitidis strains 2394 and 2169.

CLMS(15)

15. A vaccinal pharmaceutical composition which comprises, as a therapeutic agent, the lower molecular weight subunit of the human **transferrin** **receptor** of a strain of N. meningitidis, in the absence of the high molecular weight subunit of said receptor.

=> e quentin-millet, m/in

E#	FILE	FREQUENCY TERM
		HTTTTTTTTT
El	USPAT	1 QUENOUILLE, GEORGES EMILE/IN
E2	USPAT	 QUENTER, HORST/IN
E3	USPAT	0> QUENTIN MILLET, M/IN
E4	USPAT	1 QUENTIN MILLET, MARIE J/IN
E5	USPAT	2 QUENTIN MILLET, MARIE JOSE/IN
E6	USPAT	3 QUENTIN MILLET, MARIE JOSE
B/IN		
E7	USPAT	1 QUENTIN, ERIC/IN
E8	USPAT	1 QUENTIN, ERIC P F/IN
E9	USPAT	1 QUENTIN, GEORGE H/IN
E10	USPAT	1 QUENTIN, GERAD/IN
E11	USPAT	3 QUENTIN, GERARD/IN
E12	USPAT	1 QUENTIN, GERARD J/IN

- => s e4-6
 - 1 "QUENTIN MILLET, MARIE J"/IN
 - 2 "QUENTIN MILLET, MARIE JOSE"/IN
 - 3 "QUENTIN MILLET, MARIE JOSE B"/IN
- L3 6 ("QUENTIN MILLÉT, MARIE J"/IN OR "QUENTIN MILLET, MARIE JOS E"/

IN OR "QUENTIN MILLET, MARIE JOSE B"/IN)

=> d 1-6

- 1. 5,618,541, Apr. 8, 1997, Vaccine against Neisseria meningitidis infections; **Marie-Jose Quentin-Millet**, 424/250.1, 249.1; 435/871
- :IMAGE AVAILABLE:
- 2. 5,618,540, Apr. 8, 1997, Subunit vaccine against Neisseria meningitidis infections and corresponding subunits in the purified state;

- **Marie J. Quentin-Millet**, et al., 424/250.1, 249.1; 435/871 :IMAGE AVAILABLE:
- 5,045,203, Sep. 3, 1991, Separation of protein antigens of Bordetella bacteria by affinity chromatography; **Marie-Jose Quentin-Millet**,
- et al., 210/635, 198.2, 502.1, 656; 502/403; 530/413, 417, 825
- AVAILABLE:

:IMAGE

- 4. 4,985,144, Jan. 15, 1991, Affinity chromatography material for antigens of the bacteria of the Bordetella genus; **Marie-Jose B.**
 Quentin-Millet, et al., 210/198.2, 502.1, 635, 656; 502/403; 530/413,
- 417 :IMAGE AVAILABLE:
- 5. 4,965,205, Oct. 23, 1990, Culture medium for bacteria of the bordetella genus containing etherified derivative of D-glucose and a cyclodextrin; **Marie-Jose B. Quentin-Millet**, et al., 435/252, 244.
- 248, 252.1, 253.6, 822 :IMAGE AVAILABLE:
- 6. 4,774,086, Sep. 27, 1988, Process for the purification, solubilization and/or detoxification of protein antigens of bacteria of the Bordetella genus using a carbonate buffer and an acellular anti-whooping cough vaccine; **Marie-Jose B. Quentin-Millet**, et al..
- 424/240.1, 254.1, 278.1; 435/243, 244, 822; 530/417 :IMAGE AVAILABLE: